EQUALINE MUSCLE RUB- menthol, methyl salicylate cream Supervalu Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

SuperValu Inc. Muscle Rub Drug Facts

Active ingredients

Menthol 10%

Methyl salicylate 15%

Purpose

Topical analgesic

Uses

temporarily relieves the minor aches and pains of muscles and joints associated with

- simple backache
- arthritis
- strains
- bruises
- sprains

Warnings

For external use only

Do not use

- other than as directed
- on children under 12 years of age
- with a heating pad (may blister skin)

When using this product

- avoid contact with the eyes or mucous membranes
- · do not apply to wounds or damaged skin
- do not bandage tightly

Stop use and ask a doctor if

• skin redness or excessive irritation of the skin develops

- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

Keep out of reach of children

to avoid accidental poisoning. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults & children 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: consult a physician

Other information

- store at 20°-25°C (68°-77°F)
- Lot No. & Exp. Date, see crimp of tube and see box

Inactive ingredients

citric acid, glyceryl monostearate, lanolin, methylparaben, propylene glycol, propylparaben, purified water, stearic acid, trolamine

Questions or comments?

1-855-423-2630

Principal Display Panel

EQUALINE®

- non-greasy cream
- deep penetrating pain relief
- fast relief from minor arthritis, backache, muscle & joint pain

EQUALINE®

compare to Bengay® active ingredients

regular strength

muscle rub

pain relieving cream

non-greasy

NET WT 3 OZ (85 g)



EQUALINE MUSCLE RUB

Active Ingredient/Active Moiety

menthol, methyl salicylate cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-770
Route of Administration	TOPICAL		

	Ingredient Name	Basis of Strength	Strength
	MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNS PECIFIED FORM	10 g in 100 g
l	METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:0414PZ4LPZ)	METHYL SALICYLATE	15 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)		
LANOLIN (UNII: 7EV65EAW6H)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
WATER (UNII: 059QF0KO0R)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
TROLAMINE (UNII: 903K93S3TK)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:41163-770- 21	1 in 1 CARTON	01/03/2014		
1		85 g in 1 TUBE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/03/2014	

Labeler - Supervalu Inc (006961411)

Revised: 12/2021 Supervalu Inc